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(54) Title: **A PROCESS FOR PRODUCING METAL CARBOXYLATES FOR USE AS ANIMAL FEED SUPPLEMENTS**

(57) Abstract

A process for producing divalent metal carboxylates having the formula: $M(\text{CH}_3(\text{CH}_2)_x\text{COO}^-)_2$, wherein M is the divalent metal cation, zinc (Zn^{+2}) or copper (Cu^{+2}) and x is zero or 1 is disclosed. In this process, an anhydrous $\text{C}_2\text{-C}_3$ carboxylic acid is admixed with a basic divalent metal compound that is an oxide, hydroxide or carbonate of Zn^{+2} or Cu^{+2} in the absence of added solvent or other diluent. The divalent metal carboxylate so produced is used as a biologically available and economical source of trace metals for supplementation in animal diets.

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A PROCESS FOR PRODUCING METAL CARBOXYLATES
FOR USE AS ANIMAL FEED SUPPLEMENTS

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DESCRIPTION

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Technical Field

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The present invention relates to a process for producing metal carboxylates, and more particularly divalent metal acetate and propionate salts that are used for trace metal supplementation of animal feed.

Background of the Invention

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Trace elements are essential for the nutrition of animals, playing important roles in many biochemical and physiological processes. These elements include metals that form divalent cations such as zinc, copper, iron, manganese, cobalt, chromium, and molybdenum. All but molybdenum have been shown to be deficient in some natural feed ingredients, necessitating the use of supplements to make the diet nutritionally complete.

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Several chemical forms of trace metals are available for supplementation of animal diets including the inorganic salts of the trace metal, metal-amino acid complexes, metal-amino acid chelate complexes, metal-protein complexes, and metal-polysaccharide complexes (Official Publication of American Feed Control Officials, 1994, pages 209-210). U.S. Patent No. 4,315,927 also discloses the use of a metal carboxylate complex, zinc picolinate, as a food supplement for both humans and other animals. All of the complexes result

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from the complexing of a soluble metal salt with the amino acid, chelate, proteinate, polysaccharide, or carboxylic acid.

5 The salts of acetic acid, zinc acetate, manganese acetate, and cobalt acetate, have been also been approved for use as animal feed supplements (21 CFR 582.80). Zinc acetate is usually prepared in a conventional manner as is discussed below.

10 U.S. Patent No. 4,700,000 discloses that carboxylic acid salts are conventionally synthesized by reacting a carbonate, hydroxide, or oxide with a concentrated or dilute carboxylic acid. The carboxylic acid is in solution as denoted by the terms "concentrated" and "dilute". The carbonate, hydroxide,
15 or oxide are typically aqueous bases (soluble metal salts) that are reacted with the carboxylic acid to form a carboxylic acid salt. Sulfates, chlorides, and nitrates can also be used. The reactants in the conventional method are therefore in an aqueous
20 solution.

U.S. Patent No. 4,700,000 also discloses that calcium propionate is prepared by reacting propionic acid with calcium hydroxide in an aqueous solution. After concentration and crystallization, the product is
25 separated from the solution by filtration, decantation, or centrifugation, dried and ground. U.S. Patent No. 4,315,927 discloses that zinc picolinate is prepared by adding picolinic acid to an aqueous solution of a water-soluble metal salt, zinc sulfate. The product is
30 precipitated, purified by recrystallization, recovered and dried by freeze-drying.

A disadvantage of synthesis in an aqueous medium is that the use of an aqueous solution necessitates the separation of the precipitated product
35 from the solution, and drying of the recovered product.

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Another disadvantage of a conventional method is that in some cases, the divalent metal-containing base is sparingly soluble or insoluble in water.

5 Brief Summary of the Invention

 The present invention relates to an efficient and economical process for producing a divalent metal salt of a C₂-C₃ carboxylic acid; i.e., a metal carboxylate. Specifically, divalent metal acetates and
10 divalent metal propionates are contemplated, and preferably divalent metal propionates. The divalent metal C₂-C₃ carboxylates have the formula, $M(CH_3(CH_2)_xCOO^-)_2$, wherein M is the divalent metal cation, zinc (Zn⁺²) or copper (Cu⁺²) and x is zero or 1. Thus,
15 zinc acetate or propionate and copper(II) acetate or propionate are contemplated.

 The process involves agitatingly admixing anhydrous propionic acid or acetic acid with an approximately stoichiometric amount of a dry, basic
20 divalent metal compound that is an oxide, hydroxide or carbonate of Zn⁺² or Cu⁺² to form an exothermic reaction mixture. This admixture is carried out in the absence of added solvent or other diluent. The exothermic reaction mixture so formed produces water as a product
25 and a corresponding divalent metal propionate or acetate. Carbon dioxide is also formed when zinc or copper(II) carbonate are used. That exothermic reaction mixture is maintained with agitation while continually removing the formed water to form the divalent metal
30 cation propionate or acetate in dry, preferably particulate, form.

 An advantage of this invention is that it provides a process of making divalent metal carboxylates that is low in cost and easy to perform on a large
35 commercial scale. Another advantage is that preparation

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of the divalent metal carboxylate requires relatively little input of energy. Still further advantages of the invention will be apparent to a worker of ordinary skill from the description that follows.

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Detailed Description of the Invention

The present invention relates to the preparation of divalent metal propionates and metal acetates. The divalent metal C₂-C₃ carboxylates have the formula M(CH₃(CH₂)_xCOO⁻)₂, wherein M is the divalent metal cation that is zinc (Zn⁺²) or copper (Cu⁺²) and x is zero or 1. The source of the metal cation, M, is a basic metal compound that is an oxide, hydroxide or carbonate preferably zinc oxide and copper(II) carbonate.

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The following description of the present invention illustrates a process of preparing zinc propionate and copper(II) propionate using propionic acid, but the description is equally useful for the preparation of zinc acetate and copper acetate using acetic acid.

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Zinc propionate or copper propionate is prepared by admixing anhydrous propionic acid with a dry, basic divalent metal compound that is a carbonate, hydroxide, or oxide of zinc or copper(II), preferably zinc oxide or copper carbonate. The reaction unexpectedly occurs without having to dissolve the propionic acid or the basic divalent metal compound in water or another added solvent, or to suspend the reactants in any other added liquid or solid diluent. This is an advantage because the basic metal compounds used in the present invention are sparingly soluble in water.

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The propionic acid and the basic divalent metal compound are used in approximately stoichiometric

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amounts, with the propionic acid preferably being in slight excess, e.g., up to about 0.5 moles. The desired molar ratio of propionic acid to the basic divalent metal compound for the divalent cations, copper and zinc, is within the range of about 1.5:1 to about 2.5:1, preferably about 2:1, which is stoichiometric.

When used in the reaction, both the propionic acid and basic divalent metal compound are anhydrous. That is, both are substantially free of water so that neither contains more than a total of about 5 weight percent water or an aggregate of about 10 weight percent water. Preferably, the propionic acid (or acetic acid) contains less than about 0.5 weight percent water. More preferably, the propionic acid (or acetic acid) contains at most about 0.1 to about 0.2 weight percent water. The basic metal compound preferably has no waters of crystallization, as is the case with the usually reported and available forms of those basic metal compounds.

The admixed basic, divalent metal compound and propionic acid (acetic acid) are agitated together and form an exothermic reaction mixture whose components react to produce water and the copper(II) or zinc carboxylate. The exothermic reaction mixture so prepared is maintained with agitation while the water that is formed is continually removed so that the divalent metal cation propionate (acetate) that is formed is dry and free of water as discussed before.

The basic divalent metal compound is utilized in a dry, particulate form. The use of a relatively small particle size is preferred to help assure contact of the reactants and subsequent reaction. Thus, a powder form is preferred, although particles can be used that are sized to pass through at least a No. 3 sieve, U.S. Standard Sieve Series (about 1/4-inch opening),

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with no smallest sized particle being contemplated other than for the convenience of the user. Use of a high shear mixer also helps to assure that the basic metal compound particles are broken to an appropriate size.

5 Propionic acid melts at about -23° to -24°C and boils at about 142°C (about 288°F) at one atmosphere. The propionic acid is utilized in its liquid form. Propionic acid forms an azeotrope with water that boils at about 100°C (about 212°F) and
10 contains about 18 weight percent propionic acid. As a consequence of the formation of the azeotrope and the relatively low boiling point of that mixture, some propionic acid can be and is lost at higher reaction
15 temperatures as water is removed from the reaction mixture. Acetic acid melts at about 17°C and boils at about 118°C at one atmosphere. Acetic acid forms an azeotrope with water that boils at about 77°C and contains about 3 weight percent acetic acid.

 The reactants, either the $\text{C}_2\text{-C}_3$ carboxylic
20 acid or the basic metal compound or both, can be preheated relative to ambient temperature (20°C) to a temperature within the range of about 25°C to about 100°C , and preferably about 30° to about 60°C . In one preferred embodiment the $\text{C}_2\text{-C}_3$ carboxylic acid or basic
25 metal compound such as zinc oxide or both are preheated to a temperature within the range of about 30° to about 50°C , more preferably to about 40°C . However, such heating has been found to be unnecessary during large-scale production.

30 The anhydrous propionic acid is added to the dry basic metal compound in a mixing apparatus. Any reaction vessel and mixer can be used. In a laboratory procedure a beaker can be used as the mixing vessel with a stirring rod as a mixer. However, for larger scale
35 preparations, a high shear mixing apparatus is

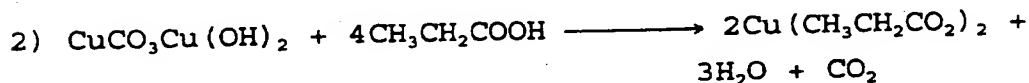
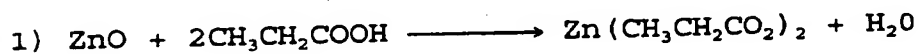
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preferred, and it is more preferred to use a high shear plow mixer. Exemplary plow mixers include the Littleford™ brand mixer (Littleford Bros. Inc.) and the Marion™ brand mixer (Marion Mixer Co., Marion, IA, Perry's Chemical Engineer's Handbook, 6th ed., Perry et al., eds., McGraw-Hill Inc., New York, 1984, page 19-18) equipped with plow-shaped and high-shear mixing blades for turbulent mixing. Additional suitable mixers can also be found in Chapter 19 of the text by Perry et al.

After thorough mixing (agitation with shear), the reaction is typically completed within minutes, but can be an hour or longer at lower temperatures as where the exotherm created by the acid base reaction is cooled.

The rate and temperature of reaction are governed by a number of variables such as the prereaction temperature of the reactants, the agitation rate within the reactor, the particle size of the basic metal compound, and the rate at which propionic acid is admixed into the reaction mixture. It is preferred to carry out the reaction at a temperature between about 150° and about 200°F (about 65° to about 93°C).

The reaction proceeds exothermically according to the following equations:



Water and heat are generated when propionic acid and the basic metal compound are reacted, as is carbon dioxide when a carbonate is used. The water along with some propionic acid is continually removed from the reaction mixture.

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In the preparation of zinc propionate, the heat of the reaction is sufficient to evaporate the water formed. In the preparation of copper propionate, additional heat is added.

5 The water is preferably released as vapor that can be aspirated under reduced pressure conditions from the exothermic reaction mixture. A well-insulated mixer is preferably employed to capture the heat given off in the exothermic reaction and evaporate water from the
10 product. The resulting product is dry and particulate.

 In one embodiment of the present invention, anhydrous propionic acid was mixed with dry zinc oxide to yield greater than 90 percent zinc propionate. The resulting zinc propionate was a white powder, and was in
15 a form readily utilizable for dietary supplementation. Unlike the conventional aqueous method of preparation, the product is not precipitated and then separated from solution. Moreover, no drying of the product by vacuum desiccation, spray drying or freeze drying is required,
20 thereby saving energy costs. In comparison, using the conventional method where the zinc oxide was dissolved in an aqueous solution of propionic acid, and the solution was heated to boiling and evaporated to yield zinc propionate crystals, the yield of zinc propionate
25 was 75.2 percent, after two crystallizations and drying under reduced pressure for 16 hours.

 The divalent metal propionate so prepared is dry (substantially free from water, as before) and particulate. It is typically and preferably a powder
30 that is itself dry but can clump slightly due to the presence of a small amount of unreacted acid. It is most preferred that the divalent metal propionate have an appearance similar to that of commercially available talcum powder. That material can be used as such, but is
35 typically recovered and packaged for sale to others.

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Recovery techniques are those normally used for non-toxic or slightly toxic powders. Purities and yields of about 80 percent and above are typically obtained, with purities and yields in excess of 90 percent being usual and preferred.

Example 1: Zinc Propionate

Two preparations of zinc propionate were made by combining 81 grams of zinc oxide and 148 grams of propionic acid in heavy-walled glass beakers. This resulted in a relative molar ratio of 1:2 of zinc cation to propionic acid. The beakers were used to insulate the reaction. One preparation utilized reactants preheated to 100°F (38°C), whereas the second preparation started at room temperature. The reactants were rapidly mixed with a stirring rod. The resulting vapor was permitted to escape from the reaction beaker. After mixing for eight minutes, a white powdery material was formed. Greater than 90 percent zinc propionate was obtained for the preheated reactants and 80 percent zinc propionate for the reactants at room temperature.

Example 2: Copper Propionate

A preparation of copper propionate is made by combining 221 grams of copper carbonate and 296 grams of propionic acid as in Example 1. This results in a relative molar ratio of 1:2 of copper(II) cation to propionic acid. The reaction is run at room temperature. The reactants are rapidly mixed with a stirring rod. The copper carbonate reacts readily with the propionic acid. Carbon dioxide is generated along with water and heat. The resulting copper propionate is a blue-green powder.

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Example 3: Zinc Acetate

5 A preparation of zinc acetate is made by combining 81.38 grams of zinc oxide and 60.05 grams of acetic acid as in Example 1. The zinc oxide reacts exothermically with the acetic acid to produce zinc acetate.

Example 4: Copper Acetate

10 A preparation of copper acetate is made by combining of 221 grams copper carbonate and 120 grams of acetic acid as in Example 1. The copper carbonate reacts exothermically with acetic acid to produce copper acetate.

15 Example 5: Zinc Propionate:Pilot Scale

Five batches of zinc propionate, 200 pounds each, were prepared using a Marion brand mixer supplied by the Marion Mixer Company, Marion, Iowa. The mixer was equipped with high-shear mixing blades set at right angles to the primary plow mixing blades. The high-shear blades were operated at a fixed speed of 3,500 rpm and the primary mixing blades were operated at 40 rpm.

25 The initial temperature of the zinc oxide and propionic acid were varied, as was the time at which the high-shear blades were engaged. Vent stack temperatures approximately six inches from the top of the mixer were monitored as a measure of the rate of reaction. Each batch was prepared by charging the mixer with 77 pounds of zinc oxide followed by spraying the dry zinc oxide with 140 pounds of propionic acid at nine pounds per minute. Starting material temperatures and the time at which the high-shear mixing blades ("choppers") were engaged are summarized below in Table 1.

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Table 1
Parameters for Pilot Scale Production
of Zinc Propionate in a High-Shear Mixer

Temperature			
Batch	Propionic Acid	Zinc Oxide	Choppers
MMZNP-1	100°F	Ambient	None
MMZNP-2	100°F	Ambient	After all propionic acid
MMZNP-3	100°F	Ambient	After 25 percent of propionic added
MMZNP-4	Ambient	100°F	After 25 percent of propionic added
MMZNP-5	100°F	100°F	After 25 percent of propionic added

Five samples, 150 g each, were taken at random from each preparation. A composite sample comprising 5 g from each sample was prepared from each preparation for analysis.

Within 48 hours of production, each pilot preparation was analyzed for yield using the above samples. Approximately ten grams of each of the above composites were weighed into a tared fritted glass funnel of 10-15 μ pore size. Sample weights were recorded to two decimal places. In each funnel, 25 mL of hexane were added and the sample stirred for one minute. The hexane was filtered off under reduced pressure, and the procedure repeated twice more. Each sample was then extracted once with 25 mL of dry acetone to remove residual hexane. Thereafter, 50 mL of water were added to each funnel and slurried with the sample for one minute. The water was then removed under reduced pressure filtration and the extraction repeated twice more for a total of 150 mL of extraction water. The residue was then extracted with 25 mL of acetone to remove residual water, then 25 mL of petroleum ether to remove excess acetone. The residue was then air-dried

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in the funnel under reduced pressure for 15 minutes. The final material was carefully removed from the funnel and weighed. Weights were recorded to two decimal places. Net conversion was calculated by subtracting the ratio of final residue weight to initial composite sample from one. Results of this analysis and physical description of pilot scale product are shown in Table 2.

Table 2
Conversion Analysis Results and Description
of Pilot Scale Zinc Propionate

<u>Batch</u>	<u>Initial Weight (g)</u>	<u>Final Weight (g)</u>	<u>Purity of ZnP*</u>	<u>Appearance</u>
MMZNP-1	10.15	1.25	87.7%	White, poorly flowing powder, hard clumps 2-20 mm
MMZNP-2	10.16	0.91	91.0%	White, poorly flowing powder, soft clumps 2-12 mm
MMZNP-3	10.18	0.64	93.7%	White, poorly flowing powder, soft clumps 2-12 mm
MMZNP-4	10.79	0.85	92.1%	White, poorly flowing powder, soft clumps 2-12 mm
MMZNP-5	10.11	0.99	90.2%	Fine, flowing white powder, minimal clumps
ZnPr-P&B ^b	10.06	0.41	95.9%	Fine, flowing white powder
ZnPr-MSF ^b	9.82	0.01	99.9%	Fine, flat white crystals
ZnO	9.95	9.89	N/A	Fine, flowing white powder

* ZnP = Zinc propionate. Based on residual solids. Not corrected for zinc oxide residual solids.

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P&B = Purchased from Pfaltz & Bauer; MSF = prepared at Kemin Industries, by reaction in dilute aqueous solution, concentration and crystallization.

5 As can be seen, purity of the zinc propionate product exceeded 90 percent in every case in which the high-shear mixing blades were used during production. The best overall purity was obtained using pre-heated propionic acid and engaging the choppers after about 25 percent of the acid had been added to the mixer. 10 However, there was little difference to be seen between this and pre-heating the zinc oxide prior to propionic acid addition. Although the lowest yield was obtained by pre-heating both reactants (MMZNP-5), this product 15 was the only one that was substantially free of clumps and free-flowing immediately after production.

When assayed by the residual solids method, zinc oxide itself showed a 0.6 percent loss in weight. Based on the appearance of the filtrate, this is due to 20 fine (<10 micron) particles and the presence of zinc hydroxide. Zinc oxide itself is soluble to only 0.00016 g/100 ml water. The zinc oxide used in this study was not less than 95 percent pure.

For the most part, stack temperatures reached 25 160°F and remained between 150°F and 200°F. In particular, MMZNP-5 reached a stack temperature of 240°F, and appeared likely to go higher.

As was noted before, propionic acid forms an azeotrope with water with a boiling point of about 30 188°F. Thus, the portions of the stack gases at stack temperatures for MMZNP-2, 3, 4 and 5 above 160°F almost certainly represent at least some azeotropic loss of propionic acid from the system along with water generated by the zinc oxide-propionic acid reaction. It 35 is believed that loss of propionic acid accounts for

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both the lower purity and dry, dusty appearance of the zinc propionate in MMZNP-5.

The reaction time was no less than 70 minutes for batches MMZNP 2-5 and 90 minutes for MMZNP-1.

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Example 6: Comparative Zinc Bioavailability

A study was conducted to compare the relative biological availability of zinc propionate, zinc methionine and zinc sulfate. The zinc preparations were added to the basal feed of day-old broilers at the levels indicated in Table 3. Each zinc treatment was replicated eight times and each pen contained ten male birds.

10

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Table 3

Design of Zinc Bioavailability Trial

Treatment	Level		Pen									
	z	ID	Source	mg/kg								
20		1	Zinc Propionate	10	13	8	21	22	35	30	49	68
		2	Zinc Propionate	20	25	20	33	16	5	18	67	56
		3	Zinpro®	10	1	14	3	34	29	12	79	86
		4	Zinpro®	20	31	2	9	46	17	48	55	74
		5	Zinc sulfate	0	37	44	27	10	23	36	91	50
25		6	Zinc sulfate	10	19	38	15	28	47	42	73	92
		7	Zinc sulfate	20	43	32	45	40	11	24	85	62
		8	Zinc sulfate	30	7	26	39	4	41	6	61	80
		9	Zinc sulfate	30	7	26	39	4	41	6	61	80

Zinpro® = Zinc methionine bisulfate; Zinpro Corporation, Minneapolis, MN.

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Basal feeds and zinc compound were mixed in a horizontal feed mixer providing the desired ingredient composition as shown in Table 4. The broilers were fed the rations for 21 days. Feed weights were taken during the trial. Body weights were measured at the beginning

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and at the end of the trial. On day 21, zinc levels in the blood and tibia ash were determined. The assay procedures for total tibia zinc content or concentration to assess zinc utilization are similar to those used by Wedekind et al., *J. Anim. Sci.*, 70:178-187 (1992). The data were analyzed according to the SAS statistical package.

Table 4
Formulation and Approximate
Analysis of Basal Feed

10	<u>Ingredient</u>	<u>Percent</u>
	Corn yellow	57.060
	Soybean meal - 48%	34.621
	Fat	3.096
15	DL methionine ¹	2.339
	Salt	0.289
	Limestone	0.621
	Defluorinated phosphate	1.724
	Vitamin premix	0.050
20	Trace min. premix ²	0.050
	Biocox ³	0.100
	Bacitracin MD-50 ⁴	0.050

¹ DL methionine is used to equalize methionine in all ratios.

² Trace mineral premix does not contain zinc.

³ Available from Hoffmann-LaRoche, Inc., Nutley, NJ.

⁴ Available from A.L. Laboratories, Ft. Lee, NJ.

Chick growth parameters and zinc levels in the blood and tibia ash were used as criteria to determine the biological availability of zinc. The broilers fed with zinc propionate had the same ending body weight as the broilers fed with zinc methionine at both the 10 mg/kg and 20 mg/kg levels. Both organic sources

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provided statistically significant higher ending body weights than did zinc sulfate at both the 10 mg/kg and 20 mg/kg levels. A similar set of results was seen for weight gain over the 21 day trial. Additionally,

5 broilers fed with zinc propionate and zinc methionine showed comparable levels of zinc in the blood and tibia ash on day 21. With these data, a relative biological availability for zinc propionate and zinc methionine can be calculated. The summary of this calculation is shown

10 in Table 5 below. The biological availability of zinc sulfate was arbitrarily set at 100%. For all parameters, there was no statistically significant difference between zinc propionate and zinc methionine with the exception of feed conversion. There was a

15 slight advantage in the broilers treated with zinc propionate compared to zinc methionine. It can thus be concluded that zinc propionate showed equal biological availability and equal performance enhancement as zinc methionine.

20

Table 5

Relative Biological Availability (%)

<u>Criterion</u>	<u>Sulfate</u>	<u>Propionate</u>	<u>Zinpro®</u>
Ending body weight (lbs)			
21 days	100.00 ^b	128.21 ^a	132.46 ^a
25 Body weight gain (lbs)			
21 days	100.00 ^b	128.24 ^a	131.52 ^a
Feed conversion (corr.)			
3-21 days	100.00 ^a	100.30 ^a	90.38 ^b
Bone ash (%) 21 days	100.00 ^b	113.18 ^a	113.95 ^a
30 Tibia ash zinc (µg/g)	100.00 ^b	125.67 ^a	123.99 ^a
Plasma Zinc (µg/ml)	100.00 ^b	113.33 ^a	124.00 ^a

Note: Means within a row grouping without a common

35 superscript are significantly different (P<0.05) as determined by Least Significant Difference.

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The foregoing description and the examples are intended as illustrative and are not to be taken as limiting. Still other variations within the spirit and scope of this invention are possible and will readily present themselves to those skilled in the art.

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Claims

1. A process for preparing a dry, particulate metal carboxylate having the formula $M(\text{CH}_3(\text{CH}_2)_x\text{COO}^-)_2$,
5 wherein M is a divalent metal cation that is zinc (Zn^{+2}) or copper(II) (Cu^{+2}) and x is zero or 1 that comprises the steps of:

(a) agitatingly admixing an anhydrous $\text{C}_2\text{-C}_3$ carboxylic acid with an approximately stoichiometric
10 amount of a dry, particulate basic metal compound that is an oxide, hydroxide or carbonate of a divalent metal cation that is Zn^{+2} or Cu^{+2} , said admixing being carried out in the absence added solvent or other diluent to form an agitated exothermic reaction mixture that
15 produces water as a product; and

(b) maintaining said exothermic reaction mixture under agitation while continually removing said water for a time period sufficient to form zinc or copper(II) $\text{C}_2\text{-C}_3$ carboxylate in dry, particulate form.
20

2. The process of claim 1 including the further step of recovering the formed zinc or copper(II) carboxylate.

25 3. The process of claim 1 wherein the basic metal compound is zinc oxide.

30 4. The process of claim 1 wherein the basic metal compound is copper carbonate.

5. The process of claim 1 wherein said $\text{C}_2\text{-C}_3$ carboxylic acid is used in excess.

35 6. The process of claim 1 wherein said $\text{C}_2\text{-C}_3$ carboxylic acid is propionic acid.

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7. The process of claim 1 wherein said C₂-C₃ carboxylic acid is acetic acid.

5 8. The process of claim 1 wherein the C₂-C₃ carboxylic acid and the basic metal compound are mixed together using a mixer with high-shear mixing blades.

10 9. The process of claim 1 wherein the C₂-C₃ carboxylic acid or the basic metal compound or both the C₂-C₃ carboxylic acid and the basic metal compound are preheated within the range of about 25° to about 100°C relative to ambient temperature prior to admixture.

15 10. The process of claim 3 wherein the C₂-C₃ carboxylic acid is propionic acid and the propionic acid or the zinc oxide or both the propionic acid and the zinc oxide are preheated within the range of about 30° to about 50°C prior to admixing.

20 11. The process of claim 10 wherein the propionic acid or the zinc oxide or both the propionic acid and the zinc oxide are preheated to about 40°C prior to admixing.

25 12. A process for preparing a dry, particulate zinc or copper(II) propionate that comprises the steps of:

30 (a) agitatingly admixing anhydrous propionic acid with an approximately stoichiometric amount of a dry, particulate basic metal compound that is an oxide, hydroxide or carbonate of zinc or copper(II), said admixing being carried in a mixer using high-shear mixing blades and in the absence of added solvent or diluent to form an agitated exothermic reaction mixture
35 that produces water as a product;

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(b) maintaining said exothermic reaction mixture under agitation while continually removing said water for a time period sufficient to form zinc propionate or copper(II) propionate in dry, particulate form; and

(c) recovering said dry, particulate zinc propionate or copper propionate.

13. The process of claim 12 wherein the basic metal compound is zinc oxide.

14. The process of claim 12 wherein the basic metal compound is copper carbonate.

15. The process of claim 12 wherein said propionic acid is used in excess.

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C07C 51/41

US CL :562/606

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 562/606

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 4,700,000 (MERKEL ET AL) 13 October 1987, col. 1, lines 6-24, col. 2 - col. 3.	1-8, 12-15
Y	US, A, 4,315,927 (EVANS) 16 February 1982, col. 1 - col. 3, line 20.	1-8, 12-15



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"A" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

02 JANUARY 1996

Date of mailing of the international search report

15 FEB 1996

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